The Vaccine Adverse Event Reporting System

VAERS ID	Vaccine Manufacturer	Adverse
909095-1	MODERNA	on 12/24/2020 the resident was sleepy and stayed in bed most of the shift. He stated he was doing okay but requested pain medication for his legs at 250PM. At 255AM on 12/25/2(
910363-1	MODERNA	Patient had mild hypotension, decreased oral intake, somnolence starting 3 days after vaccination and death 5 days after administration. He did have advanced dementia and was hc
913143-1	PFIZER\BIONTECH	Vaccine administered with no immediate adverse reaction at 11:29am. Vaccine screening questions were completed and resident was not feeling sick and temperature was 98F. At ap
913733-1	MODERNA	My grandmother died a few hours after receiving the moderna covid vaccine booster 1. While I don?t expect that the events are related, the treating hospital did not acknowledge th
914604-1	PFIZER\BIONTECH	Spouse awoke 12/20 and found spouse dead. Client was not transferred to hospital.
914621-1	MODERNA	Resident in our long term care facility who received first dose of Moderna COVID-19 Vaccine on 12/22/2020, only documented side effect was mild fatigue after receiving. She passe felt we should report the death, even though it is not believed to be related.
914690-1	PFIZER\BIONTECH	Within 24 hours of receiving the vaccine, fever and respiratory distress, and anxiety developed requiring oxygen, morphine and ativan. My Mom passed away on the evening of 12/2(
914805-1	PFIZER\BIONTECH	RESIDENT CODED AND EXPIRED
914895-1	PFIZER\BIONTECH	Injection given on 12/28/20 - no adverse events and no issues yesterday; Death today, 12/30/20, approx.. 2am today (unknown if related - Administrator marked as natural causes
914917-1	PFIZER\BIONTECH	Death by massive heart attack. Pfizer-BioNTech COVID-19 Vaccine EUA
914961-1	PFIZER\BIONTECH	pt passed away with an hour to hour and 1/2 of receiving vaccine. per nursing home staff they did not expect pt to make it many more days. pt was unresponsive in room when shol
914994-1	PFIZER\BIONTECH	pt was a nursing home pt. pt received first dose of covid vaccine. pt was monitored for 15 minutes after getting shot. staff reported that pt was 15 days post covid. Pt passed away v
915562-1	PFIZER\BIONTECH	pt received vaccine at covid clinic on 12/30 at approximately 3:30, pt vomited 4 minutes after receiving shot--dark brown vomit, staff reported pt had vomited night before. Per staff
915682-1	PFIZER\BIONTECH	Resident received vaccine per pharmacy at the facility at 5 pm. Approximately 6:45 resident found unresponsive and EMS contacted. Upon EMS arrival at facility, resident went into c
915880-1	MODERNA	Patient died within 12 hours of receiving the vaccine.
915920-1	PFIZER\BIONTECH	Resident received vaccine in am and expired that afternoon.
917117-1	MODERNA	After vaccination, patient tested positive for COVID-19. Patient was very ill and had numerous chronic health issues prior to vaccination. Facility had a number of patients who had al
917790-1	MODERNA	At the time of vaccination, there was an outbreak of residents who had already tested positive for COVID 19 at the nursing home where patient was a resident. About a week later, p vaccination caused patient's death. It simply didn't have time to save her life.
917793-1	MODERNA	Prior to the administration of the COVID 19 vaccine, the nursing home had an outbreak of COVID-19. Patient was vaccinated and about a week later she tested positive for COVID-1
918065-1	MODERNA	1/1/2020: Residents was found unresponsive. Pronounced deceased at 6:02pm
918388-1	PFIZER\BIONTECH	Resident found unresponsive without pulse, respirations at 04:30 CPR performed, expired at 04:52 by Rescue
918418-1	PFIZER\BIONTECH	Resident became SOB, congested and hypoxic requiring oxygen, respiratory treatments and suctioning. Stabilized after treatment and for the next 72 hours with oxygen saturations
918487-1	MODERNA	Two days post vaccine patient went into cardiac arrest and passed away.
918518-1	MODERNA	syncopal episode - arrested - CPR - death
919108-1	PFIZER\BIONTECH	Fever, Malaise
919537-1	MODERNA	Resident exhibited no adverse events during 30 minute monitoring following vaccine administration. Resident found without pulse at 1900.
920326-1	MODERNA	Redness and warmth with edema to right side of neck and under chin. Resident was on Hospice services and expired on 1.1.21
920368-1	MODERNA	12/30/2020 07:02 AM Resident noted to have some redness in face and respiration were fast. Resident vital signs were abnormal except blood pressure. Temp at the time was 102.(made triage aware of resident receiving Covid vaccination yesterday morning. Resident appetite and fluid consumption has been poor for few days. 12/30/2020 07:32 AM Received c as NP ordered. Will monitor for effectiveness and adverse effects if any. 12/30/2020 08:41 AM Received new orders to obtain Flu swab, obtain CBC and BMP, and Chest Xray all to be resident sent to hospital. Did educate family on benefits of Hospice services, but family persistant on continued daily care provided by nursing staff. Requests visits if decline continu Family chose at that time to sign for Hospice services and not have resident provided with IVF or IV Antibiotics
920545-1	PFIZER\BIONTECH	"The resident received is vaccine around 11:00 am and tolerated it without any difficulty or immediate adverse effects. He was at therapy from 12:36 pm until 1:22 pm when he stat call light and found he had taken himself to the bathroom. She stated that when he went to get back into the bed it was ""abnormal"" how he was getting into it so she assisted him.
920815-1	MODERNA	Found deceased in her home, unknown cause, 6 days after vaccine.
920832-1	PFIZER\BIONTECH	Vaccine 12/30/2020 Screening PCR done 12/31/2020 Symptoms 1/1/2021 COVID test result came back positive 1/2/2021 Deceased 1/4/2021
921175-1	PFIZER\BIONTECH	Resident received Covid Vaccine, noted after 30 mins with labored breathing BP 161/77, HR 116, R 38, T 101.4,
921481-1	PFIZER\BIONTECH	Vaccine given on 12/29/20 by Pharmacy. On 1/1/21, resident became lethargic and sluggish and developed a rash on forearms. He was a Hospice recipient and doctor and Hospice o
921547-1	MODERNA	DEATH ON 1/4/2021, RESIDENT RECIEVED VACCINE ON 1/2/20
921572-1	MODERNA	Resident had body aches, a low O2 sat and had chills starting on 12/30/20. He had stated that they had slightly improved. On 1/1/21 he sustained a fall with a diagnosis of a displac
921667-1	PFIZER\BIONTECH	LTCF Pfizer Vaccine clinic conducted 12/29/2020 Vaccine lead received a call indicating that a staff member deceased somewhere between 1/3/2021 and 1/4/2021. Cause of death is
921768-1	PFIZER\BIONTECH	Vaccine received at about 0900 on 01/04/2021 at her place of work, Medical Center, where she was employed as a housekeeper. About one hour after receiving the vaccine she expe hours, upon arrival to a friends home she complained of feeling hot and having difficulty breathing. She then collapsed, then when medics arrived, she was still breathing slowly ther
921880-1	PFIZER\BIONTECH	The resident was found deceased a little less than 12 hours following COVID vaccination, and he had had some changes over the last 2 days. He was 96 and had been on hospice car
922977-1	MODERNA	Fever, RespDepression & COVID positive REMDESIVIR (EUA) 200 mg x1 then 100 mg daily
923993-1	MODERNA	Patient was vaccinated Dec 30, 2020. Prime dose of Moderna vaccine. Observed for full 15 minutes post-injection. No complaints when asked during observation. Released. Subsequ home. The patient's primary care provider was unaware of his death when contacted by this reporter today (Jan 6, 2021). Electronic Medical Record without any information since th
924126-1	MODERNA	resident expired 1/1/2021
924186-1	MODERNA	Resident expired 1/3/21
924456-1	PFIZER\BIONTECH	Patient did not display any obvious signs or symptoms; the vaccination was administered at approximately 10:00 AM and the patient continued throughout her day without any comp patient's COVID-19 specimen collection from Sunday, 1/3/21, detected COVID-19. When the nursing staff went to the room to check on the resident and prepare her to move to a C
924464-1	PFIZER\BIONTECH	coughing up blood, significant hemoptysis -- > cardiac arrest. started day after vaccine but likely related to ongoing progression of lung cancer
924664-1	MODERNA	At approximately, 1855, I was alerted by caregiver, resident was not responding. Per caregiver, she was doing her rounds and found resident in bed, unresponsive, mouth open, obse incident. This PCG called 911 for EMS and gave report of incident. Resident was taken to Medical Center Emergency Department. At ER, CT scan and X-ray was performed. Per report measures. This primary caregiver reported to RN, resident recently received the first dose of COVID-19 vaccine on 1/2/21. Primary caregiver received a call from Castle RN at 0700,
925154-1	MODERNA	Deceased
925264-1	MODERNA	PT was found deceased in his home on 1/5/2021
925556-1	PFIZER\BIONTECH	Expired 1/05/2021
926269-1	PFIZER\BIONTECH	"Pt last seen at 1200 by nurse for ID band check. No visible signs of distress noted. Pt states ""I just want to be left alone"". 1230 nurse was called to pt room. Pt was noted unrespo
926462-1	PFIZER\BIONTECH	Patient developed hypoxia on 1/4/2021 and did not respond to maximal treatment and passed way on 1/5/2021
926568-1	PFIZER\BIONTECH	patient declined 12/30/2020 and was transferred to hospital where he did not respond to treatment and passed away 1/4/2020
926600-1	MODERNA	Patient did not report any signs or symptoms of adverse reaction to vaccine. Patient suffered from several comorbidities (diabetes and renal insufficiency). Patient reported not feelin
926797-1	MODERNA	had a vaccination on 12/31/2020 late morning passed away early morning 01/01/2020. This is a 93 year old with significant heart issues. EF of 20% among other comorbidities. He
927189-1	PFIZER\BIONTECH	Patient was vaccinated at 11am and was found at the facility in his room deceased at approximately 3:00pm. Nurse did not have cause of death
927260-1	MODERNA	No adverse effects noted after vaccination. Patient with cardiac history was found unresponsive at 16:45 on 1/6/21. Abnormal breathing patterns, eyes partially closed SPO2 was 41 cardiac arrest and severe bradycardia and in the end the hospital was unable to bring her back.
928062-1	PFIZER\BIONTECH	vomiting later on 01/05/21. Lethargy and hypoxia in pm of 01/06/21. Hypotension am of 01/07/21. Hospitalized, intubated, cardiac arrest, died 01/07/21.
928513-1	MODERNA	Resident passed away in her sleep
928933-1	MODERNA	Patient had been diagnosed with COVID-19 on Dec. 11th, 2020. Symptoms were thought to have started on 12/5/2020. Received Moderna vaccine on 12/23. Unexpected death on 1
929359-1	PFIZER\BIONTECH	3:07 pm lung sounds diminished oxygen sats 68%, oxygen applied Oxygen sats remained low for next 36 hours (patient on Hospice care) expired 6:22 am 1-8-21
929764-1	MODERNA	The patient was found deceased at home about 24 hours after immunization. Date of Death:: 12/29/2020; estimated time of death 6:00pm
929997-1	MODERNA	Patient received vaccine on 1/4/2021. He was in Hospice for CHF and renal failure, but was able to get up in his wheelchair and eat and take medications and talk. On 1/5/2021 am,
930154-1	MODERNA	Notified today that he passed away. No other details known at this time.
930386-1	PFIZER\BIONTECH	Patient received first dose of vaccine on 12/28, developed COVID-19 infection shortly thereafter and expired on 1/6/2021.
930418-1	PFIZER\BIONTECH	Patient received first dose of vaccine on 12/28, developed COVID-19 infection shortly thereafter and expired on 1/4/2021
930431-1	PFIZER\BIONTECH	Cardiac event, 2 days after vaccination, patient expired.
930466-1	PFIZER\BIONTECH	Fever, shortness of breath and chest pain that resulted in a heart attack a few hours after vaccination
930487-1	MODERNA	Medical docter state patient has a acute cardiac attack
930876-1	MODERNA	Death
930910-1	MODERNA	Patient received COVID vaccination around 12:15pm. Patient was monitored for the appropriate amount of time by nursing staff. Patient passed away at 2:15pm.
930912-1	PFIZER\BIONTECH	Diarrhea followed by death 24 hrs after vaccination
932346-1	PFIZER\BIONTECH	1/7-21 - Received second dose of pfizer covid-19 vaccine 1/8/21 - Fever, dizziness, headache 1/10/21 0250 was found not breathing. EMS performed CPR and patient deceased
932787-1	PFIZER\BIONTECH	RECIEVED VACCINE 1/8/21 EXPIRED UNEXPECTED 1/10/21, NO ADVERSE REACTIONS NOTED
932898-1	PFIZER\BIONTECH	The patient had an apparent cardiac arrest on 12/23/20 and was admitted to the ICU. He was taken off of life support on 12/30/20. He had known cardiac disease.
933090-1	PFIZER\BIONTECH	Patient died, I have a copy of his vaccination card
933578-1	MODERNA	Pronounced dead 1/9/2021 at 12:42. Received first dose of vaccine 1/8/2021

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933739-1	PFIZER\BIONTECH	"Staff member checked on her at 3am and patient stated that she felt like she couldn't breathe. 911 was called and taken to the hospital. While in the ambulance, patient coded. Pat determine that patient had no brain activity. Guardian, made the decision to end all life saving measures. Patient was taken off the ventilator on 1/9/2021 and passed away at 1:30a
933846-1	MODERNA	"1-2-2021 10:30 PM Complained Right arm/back hurt - took Tylenol 1-3-2021 Complained Right arm hurt, dizzy 1-4-2021 Felt better - did laundry, daughter found her deceased at 1
934050-1	MODERNA	Staff reported that patient was found Friday morning (Jan 8) sitting at a table with his head tilted forward and unresponsive to verbal or physical stimuli. Staff lowered patient to floc death of patient show that he had a fall about 1 hr. prior. It is unknown if this fall contributed to patient's death. An autopsy has been requested.
934059-1	PFIZER\BIONTECH	Acute anterior MI with death
934263-1	MODERNA	The resident resides in an independent living facility/apartment. The reporter at the center was informed by his daughter he was not feeling well on 1/1/2021 (specific symptoms cou the reporter indicates his daughter reports his test was positive.
934373-1	PFIZER\BIONTECH	Patient went to bed around 11pm on Saturday PM and sometime between then and 1:30am on Sunday morning got up and went into the living room without waking up her husband at 7:45am, she was in the recliner and did not move or anything, which is normal for her. At 8:45am, the husband went back into the living room and tried to wake his wife and that
934507-1	PFIZER\BIONTECH	Resident died suddenly and expectantly on 01/05/2021
934539-1	MODERNA	Patient received COVID-19 (Moderna) vaccine from the Health Department on afternoon of January 8, 2021 and went to sleep approximately 2300 that night. Was found unresponsiv
934963-1	PFIZER\BIONTECH	Death; This is a spontaneous report from a contactable Physician. An elderly male patient received BNT162B2 (COVID vaccine), via an unspecified route of administration on an unsp was unknown if an autopsy was performed. It was unknown if any treatment was received for the event. It was unknown if the patient was diagnosed with COVID prior vaccination o vaccine] in the country of incidence or the country where the product was purchased (if different). This may be a duplicate report if another marketing authorization holder of [COVII assessment. Further information such medical history, concomitant medications, concurrent illness and event term details especially death cause and autopsy results are needed for aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorit
934966-1	PFIZER\BIONTECH	COVID-19; COVID-19; Pneumonia; respiratory failure; This is a spontaneous report from a contactable consumer. An 80-year-old female patient received first dose of BNT162B2 (PF Concomitant medications included unspecified medications. The reporter's mother in law was tested for COVID-19 at a nursing facility on 25Dec2020 and she was negative. On 02Jar death was listed as pneumonia, respiratory failure and COVID-19. No autopsy performed. No treatment received. No one knew if the vaccination contributed to her death. It was har reporter wanted to give us this information because we might want to consider having high risk population, patients with underlying conditions, older population tested for COVID-19 did not receive any other vaccines within 4 weeks prior to the COVID vaccine. There are medications the patient received within 2 weeks of vaccination. Prior to vaccination, the patie Sender's Comments: The association between the fatal event lack of effect (pneumonia, respiratory failure and COVID-19) with BNT162b2 can not be fully excluded. The impact of th safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators Pneumonia, respiratory failure and COVID-19
934968-1	PFIZER\BIONTECH	he passed away; not responsive; mind just seemed like it was racing; body was hyper dried; Restless; not feeling well; ate a bit but not much; kind of pale; Agitated; Vomiting; trou an unspecified route of administration, on 04Jan2021 (at the age of 54-years-old) as a single dose for COVID-19 immunization. Medical history included diabetes and high blood pres UNKNOWN), and amlodipine (MANUFACTURER UNKNOWN). The patient experienced not feeling well, ate a bit but not much, kind of pale, vomiting, trouble in breathing, and agitatec as: around 10:15 AM). The clinical course was reported as follows: The patient received the vaccine on 04Jan2021, after which he started not feeling well. He went right home and w Emergency services were called, and they took his vitals and said that everything was okay, but he was very agitated; reported as not like this prior to the vaccine. The patient was t just dried up. The patient continued to vomit throughout the day and then he was very agitated again and would fall asleep for may be 15-20 minutes. When the patient woke up, he morning, he was still agitated. The patient stated that he couldn't breathe, and his mind was racing. The patient's other brother went to him and he was not responsive, and he pass aforementioned. The clinical outcome of all of the events was unknown; not responsive was not recovered, the patient died on 06Jan2021. The cause of death was unknown (reporte Death: not responsive and he passed away
935222-1	MODERNA	Patient was reported to be deceased at home by law enforcement on 1/7/21
935343-1	PFIZER\BIONTECH	There were no adverse reactions. Resident Died, she had a history of issues with her health prior to the vaccine.
935350-1	MODERNA	Patient was found unresponsive at home with SpO2 20% 1/2/2021
935511-1	MODERNA	Patient received the 1st dose of Moderna and was found deceased in her home the next day.
935767-1	PFIZER\BIONTECH	My mother was given Pfizer vaccine on Thursday and she died 3 days later yesterday on Sunday!!!
935815-1	PFIZER\BIONTECH	Difficulty breathing, death.
936043-1	MODERNA	RESIDENT 1ST DOSE OF MODERNA VACCINE ADMINISTERED ON 01/04/2021 AT 8:30PM, RESIDENT FOUND UNRESPONSIVE ON 01/05/2021.
936738-1	PFIZER\BIONTECH	loss of consciousness Narrative: Patient received COVID-19 vaccine dose #1 on 1/6/21 w/o complications. Per 1/6/21- 1/9/21 nursing notes, patient did not experience any injection bed. Per nurses, he was previously awake/alert, talking and asymptomatic. Patient is DNR/DNI but facility rapid response emergency team called d/t patient's sudden change of conc breathing. Pulse ox 94%, HR in 60s per machine. BP unmeasurably low by BP cuffx3. Resident passed at 18:20 pm.
936805-1	MODERNA	Patient received the vaccine on 12/22/20 without complication. It was reported today that the patient was found unresponsive and subsequently expired at home on 1/11/21.
937127-1	MODERNA	The facility had positive cases of COVID when we were able to begin vaccinating residents. Within about a week of vaccination, patient was tested positive for COVID. He was 91 yea
937152-1	MODERNA	The facility had positive cases for COVID 19 when the vaccine was received and administered to patient. With her advanced age and chronic conditions, she did not have time to buil
937186-1	MODERNA	The facility had a number of positive COVID 19 cases prior to patients vaccination. Due to her advanced age, chronic condition, and exposure, patient did not have the time to build i
937434-1	MODERNA	Pt expired due to possible cardiac arrest. Unsure if this was vaccine related.
937444-1	PFIZER\BIONTECH	Resident was found deceased at approximately 6pm in her apartment
937527-1	PFIZER\BIONTECH	unsure if related to vaccine, but was notified by her next of kin that she died on 1/4/2021. No reports of side effects or hospitalization were reported to the facility prior to the notific
937569-1	MODERNA	patient reported expired 1/7/2021
937773-1	PFIZER\BIONTECH	Patient was sent to the ED due to significant hematuria. He was afebrile.
938097-1	PFIZER\BIONTECH	died; This is a spontaneous report from a non-contactable consumer via a Pfizer-sponsored program. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH reported. It was reported the patient was a doctor, died after the vaccine with no apparent disease. It was not reported if an autopsy was performed. No follow-up attempts are poss
938118-1	PFIZER\BIONTECH	on 1/8/2021 17:30 patient taken to ER, cerebellar hemorrhage, stroke, aneurysm
938974-1	PFIZER\BIONTECH	Hospice Resident received first Covid 19 vaccine dose on 1/6/21. 1/7/21 resident had decreased appetite noted in am but ate 100% of meal at dinner. 1/9/21 resident had decreasec resident not eating meals but ingesting milkshake and medications without any problems. Hospice contacted for change in condition. 1:00 pm hospice ordered Phenergan 12.5 mg Q
939050-1	MODERNA	Patient vaccinated on 12/28. Approximately one day later, develops cough and on azithromycin x 1 week. On 1/3, patient develops left-sided weakness and aphasia. Taken to the hos
939270-1	PFIZER\BIONTECH	Sudden cardiac death
939845-1	PFIZER\BIONTECH	Three hours after receiving COVID 19 vaccination, Patient oxygen level decreased to a critical level and went into cardiac arrest. Staff performed full code but was unable to bring ba
940602-1	MODERNA	"Patient received vaccine on 1/8/2021. On 1/9/2021 I checked on patient via phone for symptoms or problems and he reported none but mild soreness at injection site. On 1/10/20 appeared to be a ""heart attack""."
940822-1	PFIZER\BIONTECH	patient passed away after receiving the Covid vaccine; This is a spontaneous report from a contactable nurse. An 81-year-old male patient received BNT162B2 (PFIZER-BIONTECH C 08Jan2021, the patient passed away after receiving the COVID vaccine. The patient died on 08Jan2021. An autopsy was not performed. Investigations indicate that unspecified labs thought that he potentially passed away from the COVID vaccine. The relatedness of the event to the suspect vaccine was reported as related by the reporting nurse per The Agency possible to make meaningful causality assessment, it is unlikely the vaccine could have contributed to the death of the patient based on the known safety profile. However case will b safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in res the Covid vaccine
940855-1	MODERNA	Patient received her vaccination on 1/12/21 administered by pharmacy*+. She expired on 1/12/21 an approximately 7:30pm. Resident did not have any adverse reactions and was
940866-1	MODERNA	"Patient was found ""acting abnormal"" on 1/9/2021 at 1215. VS HR 20-30's. EMS activated. EMS arrived and patient was found pulseless in PEA/ asystole, CPR and ACLS initiated a
940950-1	PFIZER\BIONTECH	thrombopenia; pulmonary embolism; neutropenia fever; This is a spontaneous report from a Pfizer-sponsored program . A contactable consumer reported for a patient that received history and concomitant medications were not reported. The patient experienced thrombopenia, pulmonary embolism and neutropenia fever on an unspecified date. The clinical outc vaccine, BNT162B2, was not provided and will be requested during follow-up.; Reported Cause(s) of Death: thrombopenia; pulmonary embolism; neutropenia fever
940954-1	PFIZER\BIONTECH	"Heart attack; This is a spontaneous report from a contactable consumer. An 82-year-old female patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VAC doctor's office/urgent care. The patient's medical history and concomitant medications were not reported. It was unknown if the patient received any other vaccines within four week patient experienced heart attack; which resulted in death and was assessed as medically significant. The patient also experienced the associated symptoms of cold sweats, chest pai outcome of the event, heart attack, was fatal. The patient died on 05Jan2021 due to heart attack; as ruled by the paramedics. It was unknown if an autopsy was performed. The bat
940955-1	PFIZER\BIONTECH	"Cardiac Arrest; Patient was found pulseless and breathless 20 minutes following the vaccine administration.; Patient was found pulseless and breathless 20 minutes following the va second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL1284) via intramuscular at left arm on 11Jan2021 12:15 PM at single dose for COVID-19 immuniza patient received medication within 2 weeks of vaccination included amiodarone, melatonin, venlafaxine hydrochloride (EFFEXOR), ibuprofen, aripiprazole (ABILIFY), lisinopril, cranbe alexandrina leaf (SENNA [SENNA ALEXANDRINA LEAF]), polyethylene glycol 3350 and morphine. The patient did not receive any other vaccines within 4 weeks prior to the COVID va right arm on 21Dec2020 12:00 PM at single dose for COVID-19 immunization. Since the vaccination, the patient been tested for COVID-19 (Sars-cov-2 PCR) via nasal swab on 06Jar Patient died on 11Jan2021 12:30 AM because of cardiac arrest. No treatment received for the events. Outcome of pulseless and breathless was unknown. the autopsy was performec disabling/Incapacitating nor congenital anomaly/birth defect.; Sender's Comments: Based on the available information this patient had multiple underlying medical conditions includi death. However, based on a close temporal association (""Patient was found pulseless and breathless 20 minutes following the second dose of BNT162B2 vaccine administration, con Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any ap determined Cause(s) of Death: autopsy remarks was unknown. Autopsy-determined cause of death was unknown"
941215-1	PFIZER\BIONTECH	Actual event and cause of death were unknown; This is a spontaneous report from a non-contactable consumer. A 90-year-old female patient received first dose of BNT162B2 (PFIZE from Nov2019. Concomitant medications were not reported. The consumer stated that she was taking the reporting responsibilities to report that a friend of hers, informed that the herself, regarding the incident. Their conversation was very brief. The patient was 90 years old, and it was her friend's mother that was the patient. Actual event and cause of death autopsy was unknown. The outcome of the event was fatal. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Reported Cause(s) of Deat
941561-1	MODERNA	Staff walked into resident's room around 10:00am and noted resident's left side of his face was flaccid. Nurse was called and upon assessment resident noted to have an unequal ha ability to use his left side. Resident passed away on 1/11/2020.
941607-1	MODERNA	The patient passed away today, 1/13/2021. She was a hospice patient. She showed no adverse effects after receiving the vaccine on 1/12/2021. This morning she woke up as norma
941743-1	MODERNA	This person was found to be deceased on routine rounds during the night, 3am. No symptoms of reaction noted post vaccine. No injection site reaction. No reports of any allergic rea
941811-1	MODERNA	Resident began having fever on 1/11/21 @0600. VS= T-102 B/P- 100/57 P- 112 RR- 24 O2 Sat 92% on RA. MD called. Rapid COVID Test was negative. CBC,CMP, U/A were ordered ordered. Condition worsened. Resident sent to nearest ER for evaluation. Later in the evening the staff AT Medical Center called to inform staff that resident had expired @ 2230 as a
942040-1	PFIZER\BIONTECH	little bit of a reaction light headed after 5 minutes. vitals were low, so observed for 30 minutes after being light headed. Patient was found unresponsive and pronounced dead later t
942072-1	PFIZER\BIONTECH	Death occurred 3 days after vaccine receipt; attributed to complications of her chronic advanced dementia with aspiration at age 87. No evidence of acute vaccine reaction.
942085-1	PFIZER\BIONTECH	No adverse effects from vaccination seen on 1/2/21. On 1/6/21 resident was seen by Dr and her baclofen pump was refilled with 20 ml Baclofen 4,000mcg/ml. ITB Rate increased by primary physician was notified as well as her daughter. Oxygen increased to 4 L/min, sats at 83%. SOA noted, reported all over pain. At 0850 when they attempted to reposition the
942106-1	PFIZER\BIONTECH	54 y/o M with PMH of HTN, HLD, Alcoholic Cirrhosis, Aortic Valve Stenosis, and angina BIBA as a Medical Alert for cardiac arrest noted PTA. Per EMS, the patient called because he wa had no associated shockable rhythm. Of note, pt's wife, had noted pt had received covid vaccine the prior day.
942290-1	PFIZER\BIONTECH	Resident received 1st dose on 1/4/2021. On 1/6/2021 resident having SOB, increased weakness with O2 sats at 91% RA. On 8th resident sustained a fall, O2 sats 88-92, dizzy, wea from the ER on 1/9/2021 with new diagnosis of Leukemia and orders for hospice. Continued with fever, crackles and N/V and loss of appetite from the 9th and 10th of January. Resid
943266-1	PFIZER\BIONTECH	Initial pain in back of head and extreme headache. Some vomiting. At emergency, went into coma and was intubated. Hole drilled in skull to relieve pressure. MRI taken. Lot of bleec
943362-1	MODERNA	Pt collapsed at home approx 5:30 pm and died
943397-1	PFIZER\BIONTECH	On day due for 2nd dose, Patient was found unresponsive at work in the hospital. Patient pupils were fixed and dilated. Full ACLS was initiated for 55 minutes with multiple rounds of

VAERS ID	Vaccine Manufacturer	Adverse
943442-1	PFIZER\BIONTECH	Systemic: reported by staff patient expired under suspicious circumstnces after receiving vaccine. Patient was on hospice, reported not expected to pass this soon; symptoms lasted
943889-1	MODERNA	No adverse reactions observed after administration of medication. Patient starting complaining of shortness of breath around 0500 the following morning. SP02 checked in the 80s. F
944282-1	PFIZER\BIONTECH	resident coded on 09Jan at 8am and expired; This is a spontaneous report from a contactable Other Health Professional. A 70-year-old male patient received first dose of BNT162B2 DM2(Type two diabetes mellitus), CHF(congestive heart failure), open wound, wound infection, heart failure. Allergies to medications, food, or other products: none. Concomitant me the COVID vaccine: Unknown. Facility where the most recent COVID-19 vaccine was administered: Nursing Home/Senior Living Facility. The resident coded on 09Jan2021 at 8 AM an Unknown. Prior to vaccination, was the patient diagnosed with COVID-19: No. Since the vaccination, has the patient been tested for COVID-19: No. Serious: Yes. Seriousness criteria Congenital anomaly/birth defect: No.; Sender's Comments: The old patient had diabetes mellitus, congestive heart failure, open wound complicated by infection, all these pre-existir results are needed for a full assessment of the case. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evalu notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate; Reported Cause(s) of Death: resident coded on 09Jan at 8am and expired
944365-1	PFIZER\BIONTECH	Resident expired on 12/30/20, dx cardiac arrest.
944439-1	PFIZER\BIONTECH	Resident expired on 1/2/21.
944595-1	PFIZER\BIONTECH	Cardiac arrest within 1 hour Patient had the second vaccine approximately 2 pm on Tuesday Jan 12th He works at the extended care community and was in good health that morning hour of the 2nd vaccine. EMS called immediately worked on him 30 minutes in field then 30 minutes at ER was able to put him on life support yet deemed Brain dead 1-14-21 and pl
944641-1	MODERNA	Patient died on 1/21-2021
944732-1	MODERNA	Resident found unresponsive and without pulse at 05:45am.
944998-1	PFIZER\BIONTECH	On 1/11/21 noted with headache, nausea/vomiting, severe melaise. On 1/12/21 resident expired.
945241-1	PFIZER\BIONTECH	71yo female resident who died after receiving Pfizer BioNTech vaccine. On 1/14/2021, VS taken at 10am, B/P 99/60, O2 sats, 95% (trach w/O2). At 11:30am, Patient showed no s/s was immediately started; no shock advised per AED; 12:15pm EMS arrived and took over. At 12:38pm, EMT called time of death.
945247-1	PFIZER\BIONTECH	Has underlying dementia and often with difficulty eating. 1 week after immunization she developed a stroke with left sided weakness and difficulty swallowing. Comfort measures ins
945253-1	PFIZER\BIONTECH	"83yo female resident who died after receiving Pfizer BioNTech vaccine. On 1/14/2021, the patient reportedly got up in the middle of the night with c/o feeling ""blah"", restlessness, unresponsive, no pulse, no respirations. GNA notified the nurse. At 6:03am, CPR started and EMS called. At 6:15am, EMS arrived and took over. At or around 6:30am, EMT called tin
945578-1	PFIZER\BIONTECH	No reactions immediately after vaccine was given. Resident has dementia, has had multiple hospitalizations related to a renal stone recently. Had a tooth that was bothering her, wer eating. Compassionate visit with family. Family did not want hospice, did not feel it was needed, said, what more could they do for her than you're already doing? On 1/11 at 1950 w
945603-1	PFIZER\BIONTECH	Had no immediate issues with the vaccine. He had returned from the hospital on 12/21 and had some concerns about his weight which were shared with his physician on 1/4/21. On
946225-1	PFIZER\BIONTECH	At approximately 10:30pm on 1/14/2021, resident was noted to have a rash on her face, hands, arms, and chest. VS:100.2, 113, 20,108/59, 84% room air. applied nasal cannula at 1/15/2021, resident congested and coughing. BP 151/70, pulse 124, temp 98.1 forehead, resp 20 and pulse oc 79% on 3L. At approximately 2:30am PRN cough syrup and breathing called to say resident passed away.
946293-1	MODERNA	51 year old M with h/o O2 dependent COPD, Severe pulmonary fibrosis became increasingly hypoxic around 1800hours 1/7/2021. He was transported to hospital for acute on chroni medical center.
946959-1	PFIZER\BIONTECH	Sudden death 18 hours post vaccine .
947129-1	MODERNA	Resident received Moderna vaccine on 12/23/2020 around 5 pm. At approximately 3:35 am on 12/25/2020, resident had a CVA and died on 1/1/2021 at 3:00 am.
947642-1	PFIZER\BIONTECH	died two days after receiving the vaccine; Fever; This is a spontaneous report from a contactable consumer (patient's stepchild). A 66-year-old male patient received the second dos The patient's medical history was not reported. Concomitant medications included an unspecified statin. The patient experienced fever on 08Jan2021. The patient died two days afte the middle of night. It was reported that it was not clear what exactly happened, but they are looking into this. The clinical outcome of fever was unknown and of died two days afte number for the vaccine, BNT162B2, was not provided and has been requested during follow up.; Reported Cause(s) of Death: died two days after receiving the vaccine
947662-1	MODERNA	Accelerated decline in condition with decreased input, decreased responsiveness, somnolence, and death
947841-1	MODERNA	Patient had no immediate effects from the vaccine, but died approximately 8 hours after receiving first dose of vaccine.
947974-1	PFIZER\BIONTECH	Resident was found without a pulse and not breathing 20 minutes after vaccine administration. Upon MD review, no signs of anaphylaxis were noted.
948150-1	PFIZER\BIONTECH	increase weakness and fatigue, weakness in extremities, incontinent, jerky arm movements, within first 24 hours, continue to decline sent to hospital returned weaker, within 24 hrs
948164-1	MODERNA	Abdominal pain, Headaches, chest pain, loss of appetite, confusion, elevated liver enzymes 1/8-1/15/21
948181-1	MODERNA	Death Chest pain; irreg heart rhythm; evening of vaccine; death on toilet on 1/13/21
948228-1	PFIZER\BIONTECH	Patient reportedly expired the day following receipt of the vaccine.
948418-1	PFIZER\BIONTECH	Expired on 1/12/2021; unknown cause of death
948428-1	MODERNA	We got a call from a home health nurse Brandu Talamo, stating that the patient passed away.
949474-1	MODERNA	Resident had lunch on 01/14/21 and after lunch around 2:00pm, he vomited and stopped breathing. We coded the resident and 911 paramedics came. They pronounced him dead at
949523-1	MODERNA	Around 00:50am on 01/15/21, C.N.A. reported that the resident looked different and not responding. Initiated Code Blue and started CPR. 911 arrived and pronounced resident deac
949547-1	PFIZER\BIONTECH	"The patient stated "" I just feel Blah"". vital signs obtained. 156/75 p-84 spo2 94% via NC 2L. T-96.7, c/o feeling restless, c/o nausea with no vomiting. Patient observed at 0600 nc
949630-1	MODERNA	This patient has been under hospice care for over 2 years at the nursing home. She has had a steady decline with gradual weight loss. She was totally dependent in her care needs. hours for first 72 hours after vaccination) with BP 64/52 but otherwise asymptomatic. Subsequent BP improved. On 1/4/2021 at 4:45 am, pt found with respiratory rate of 30 with o after that point had persistent tachypnea and worsening hypoxemia despite clear lungs on exam. She remained under hospice care and comfort measures were continued. No blood
949657-1	PFIZER\BIONTECH	Veteran was found by family slumped over and unresponsive at the breakfast table on 1/13/21, had expired
949965-1	PFIZER\BIONTECH	Patient 101 years old, nursing home resident, received vaccine 1/11, on 1/13 found on floor without obvious trauma, unresponsive. Brought to ED and was bradycardic, hypotensive, accepted case although initially unknown that patient had recently received vaccine. ME updated with that information today as soon as discovered.
950057-1	MODERNA	Patient suffered a cardiac arrest and was unable to give details about her symptoms. Per husband, patient did not complain of any symptoms after vaccine administration. She begar
950073-1	MODERNA	"On 1/15/2021 at 1800, resident noted to be lethargic and shaking, stating ""I don't care."" repeatedly. C/O head and neck pain. T100.6. Given Tylenol with no relief of pain. Order r , T 99.4, Absence of vital signs at 4:15AM 1/16/21 and death pronounced at 4:40AM 1/16/21."
950108-1	MODERNA	"""Moderna COVID-19 Vaccine EUA"" It has been reported to me that pt. had gone into hospital for a heart catheterization on 1/12/2021. It was found during this procedure that pt.
950441-1	PFIZER\BIONTECH	Pt had witnessed arrest by wife. Pt wife started CPR and called EMS. CPR started at 15:12. Continued by EMS. Pt arrived to medical center asystole with CRP in progress and ventilat unresponsive to verbal and tactile stimulus and had fixed unreactive pupils. He was pronounced at 16:13.
950893-1	PFIZER\BIONTECH	Death
950935-1	PFIZER\BIONTECH	Resident expired
950979-1	MODERNA	Headache after dose was given at 10:00 a.m Died at after 7:30 pm the same night the dose was given.
951101-1	PFIZER\BIONTECH	PATIENT GOT HER FIRST COVID PFIZER VACCINE AT 12/31 IN THE AM. HAD GOTTEN FLU LIKE SYMPTOMS AND HAD BEEN SICK FOR A COUPLE OF DAYS. HAD NAUSEA AND VOMIT STROKE EXAM. PT HAD NO MOVEMNET IN ARMS OR LEGS AND WAS UNABLE TO SPEAK. PT WAS VITALLY STABLE AT THE TIME. EMS RECORDED THAT THEY THOUGHT DIAGNOSIS
951518-1	MODERNA	"Narrative: Patient with severe aphasia and only able to say ""hey, hey, hey"" or ""uh huh"" or shake his head no as a way to communicate. Patient previously able to ambulate with aggressive behavior of shouting ""hey"" and grabbing of groin in 2016. This was worked up with CT scans, labs, referral to urology, neurology, and referrals to psychiatry. The exact reduced, and improved again with addition of injectable antipsychotic on 12-10-2020.Patient suffered from falls on occasion given his significantly impaired physical mobility. His last often refuse medications. He would sometimes indicate that they would cause dizziness, and other times he would simply refuse. We attempted to hide medications in his food/fluid clear indication to continue use. He was high fall risk and would often refuse this medication as well since 10/2020. Noted to be in NSR on EKGs and decision made to discontinue the change in health condition. Temperature 36.8Con January 4th at 19:45. During routine bedtime cares, patient suddenly collapsed and death was pronounced January 4, 2021 at 20:
951519-1	PFIZER\BIONTECH	Narrative: Symptoms: Palpitations & Syncope Treatment: EPINEPHRINE 1 MG ONCE ,EPINEPHRINE 1 MG ONCE ,SODIUM BICARBONATE 50 ML ONCE
951678-1	PFIZER\BIONTECH	Heart attack death medical test
951688-1	MODERNA	Resident expired 1/17/21
952204-1	PFIZER\BIONTECH	Patient became sick 3 hours after the vaccine and was found deceased 1 day after his vaccination. He passed away in his sleep.
952704-1	PFIZER\BIONTECH	Daughter call in for VAERS report to file for father whom committed suicide 1/16/2021 in the AM after reportable ae of COVID 19 vaccine administered 1/14/2021. Patient sought ca summary diagnosis: adverse reaction to COVID shot, fever, Panic Disorder-- ER. Medical Center Discharge summary diagnosis: Adverse reaction to the vaccine, acute anxiety. Report would get up and sit down again repeatedly, agitated and anxious. Attempted to urinated hospital bed. Patient committed suicide in home.
952713-1	MODERNA	Weakness, Low O2, death. Positive for COVID on 1/12/21, dies on 1/16/21
952799-1	PFIZER\BIONTECH	On 1/17/2021 at 4:35 am resident found apneic and pulseless, at 4:40am death confirmed
952881-1	MODERNA	Resident was seen by MD on 1/11/2021 due to increasing in edema and shortness of breath. Lasix 40 mg STAT given. New orders to get a STAT CBC, CMP, and BNP. Resident has be was assisted to the toilet on 1/15/2021 in the morning where he subsequently passed away.
953129-1	MODERNA	Patient presented to our Emergency Department via EMS in full code status; asystole. Patient expired. Per nursing, husband stated patient awoke this AM and reported pain in back t
953183-1	PFIZER\BIONTECH	1/11/21 at 8:57 Resident with fever and at 11 am saturation down to 83 O2 to 10 liters. Resident continued to decline until CTB on 1/14/2021 at 1325
953348-1	MODERNA	Patient was living in a nursing home with positive cases when administered. His age and chronic condition was such that he did not have time after the vaccination to avoid exposure
953590-1	PFIZER\BIONTECH	resident expired; This is a spontaneous report from a contactable healthcare professional. An 82-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVI from, failure to thrive (FTT), diabetes mellitus (DM) 2 , chronic obstructive pulmonary disease (COPD), arthritis, weakness, hyperlipidemia, chronic kidney disease (CKD), dementia. negative on 09Jan2021. There was no treatment given for the event. The patient died on 11Jan2021. An autopsy was not performed.; Sender's Comments: Lacking information on th measure and for reporting purposes. The patient's pre-existing medical condition of metabolic encephalopathy from, failure to thrive (FTT), diabetes mellitus (DM) 2 , chronic obstruc impacts of this report on the benefit/risk profile of the product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for Investigators, as appropriate.; Reported Cause(s) of Death: resident expired
953754-1	PFIZER\BIONTECH	patient suddenly developed pneumonia 7 days after vaccination and died the evening of developing pneumonia
953785-1	MODERNA	Death
953858-1	MODERNA	patient started to decline 1/10/2021, patient seen at facility by medical professional - patient deceased 1/13/2021
953865-1	PFIZER\BIONTECH	REPORTING ONLY AS RESIDENT EXPIRED ON 1/17/2021 3 DAYS AFTER. S/S HYPOXIA/CONGESTED LUNG SOUNDS
953922-1	PFIZER\BIONTECH	The day following the vaccine, the patient complained of throat issues and anxiety. This was not new... however . That evening he reported difficulty breathing and was placed on ox obtained from the hospital indicated the patient died from a massive myocardial infarction.
954251-1	PFIZER\BIONTECH	71 year old woman at rehabilitation center for physical therapy with history of cirrhosis of the liver, asthma, and heart condition was tested for COVID-19 on 01/07/21, received 1st confused state. Patient passed away on 01/17/21.
954780-1	MODERNA	On 1/13/2021, resident had sudden emesis. Immediately following emesis he was noted without a pulse and pronounced deceased. No acute symptoms noted prior to this episode.
954812-1	PFIZER\BIONTECH	She had the first dose of Pfizer vaccine at the Campus on Friday 1/15 at 4:30 pm. After the vaccine, she had no new symptoms or signs of vaccine reaction and MD friend reports tha nausea/epigastric pain, and chest heaviness. These apparently were not unusual symptoms for her to feel intermittently. Per her niece, who has a home O2 sat device, her 02 sat tha
955256-1	PFIZER\BIONTECH	Patient was vaccinated in right arm. Within 5 to 10 seconds after vaccination, patient started clinching his hands tightly and became unresponsive. Patient was lowered to the floor a
955261-1	PFIZER\BIONTECH	Death

VAERS ID	Vaccine Manufacturer	Adverse
955390-1	PFIZER\BIONTECH	Resident received vaccination on January 15, 2021. She was found unresponsive with shallow respirations on the morning of January 16, 2021 and was sent to ER via ambulance. Th
955425-1	MODERNA	resident had a pressure ulcer to RT hip, was getting treatment on. Was scheduled to have wound debrided and wound vac applied on 1-19-2021. Appetite was poor, not wanting to g
955436-1	PFIZER\BIONTECH	patient received vaccine 12/29. Unexpected death 1/5.
955532-1	PFIZER\BIONTECH	COVID 19 Vaccination administered by pharmacy staff. No adverse effect at the present time. Staff will continue to observe adverse reaction. Will continue to monitor. Patient at star floor 911 arrived at the scene at 3:10am Cpr rotated Between Nursing and EMT on Scene. Cpr was given to patient for over 45 minutes. Patient was pronounced at the scene at 3:5(commode unresponsive with absent respiration and pulse. Resident lowered down on the floor with 4 person assist. CPR initiated, AED pads placed on chest with no shock indicated. for family to call us back for funeral arrangements.
955597-1	PFIZER\BIONTECH	Death
955878-1	PFIZER\BIONTECH	his platelet levels dropped and he had a hemorrhagic stroke; his platelet levels dropped and he had a hemorrhagic stroke; This is a spontaneous report from a contactable consumer immunization. The relevant medical history and concomitant medications were not reported. The patient died 2 weeks after his COVID shot because his platelet levels dropped and h Reported Cause(s) of Death: his platelet levels dropped and he had a hemorrhagic stroke; his platelet levels dropped and he had a hemorrhagic stroke
955879-1	PFIZER\BIONTECH	expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar death events for 8 patients. This report is for 5th of 8 pa dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecifie information provided in this report does not allow a full assessment of the case. The event death with unknown cause is assessed as related to the suspect drug per company guidan available.,Linked Report(s) : US-PFIZER INC-2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: expired before receiving the second dose
955880-1	PFIZER\BIONTECH	passed unexpectedly; This is a spontaneous report from a contactable nurse communicated to a Pfizer colleague. This nurse reported similar death events for 8 patients. This report dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient passed unexpectedly on an unspecified date. The patient report does not allow a full assessment of the case. The event death with unknown cause is assessed as related to the suspect drug per company guidance. This case will be reasses reassessed if additional information becomes available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety ev promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.,Linked Report(s) : US-PFIZER INC-2021034595 same drug, reporter and event but
955959-1	MODERNA	Patient died 1 week after vaccination. According to family was having very rapid decline in status in recent weeks and they did not think related to vaccination.
956225-1	PFIZER\BIONTECH	Systemic: Pt monitored by nursing for 30min after inj,pt was stable/no reaction.At ~1hr post inj pt was unresponsive.Pt was a hospice/dnr per director
956365-1	PFIZER\BIONTECH	12/28/2020: generalized weakness and fell twice at home, cough, nausea,1/04/2021: cough, nausea, fever and chronic pain when she fell from being weak. admitted to hospital wit passed away
956458-1	PFIZER\BIONTECH	Patient was vaccinated for SARS-CoV-2 on 6-Jan-21 at his site of employment, a Nursing Home. Patient presented to Urgent Care on 15-Jan-21 complaining of left sided chest pain t a slightly prolonged QTc of 463 ms. Physical exam was significant for bibasilar crackles and X-ray showed bibasilar infiltrates consistent with COVID pneumonia but bacterial pneumo guaifenessin with codeine cough syrup, and Zofran. Labs were drawn and he was discharged. His lab results were reported after his departure and were significant for a white blood alkaline phosphatase of 294 and AST of 112 with ALT noted to be within normal limit. His COVID nasopharyngeal swab from the visit was reported as negative and a swab performec Department received a 911 call about an adult at the patient's address who was found unresponsive. Upon arrival on scene, the patient was found to be deceased and a decision was Examiner's Office. They have decided to perform an autopsy and have recovered the CBC and chemistry specimens obtained for further testing.
956761-1	PFIZER\BIONTECH	Family was told that Patient expired in his sleep during the early morning hours of 1/15. I spoke with him the evening before (on 1/14), which was a day after he had received the C
956811-1	MODERNA	Resident was noted unresponsive, no respiration, no blood pressure, no pulse, code blue called according to facility protocol, resident is full code, CPR started, 911 called, arrived an
956843-1	PFIZER\BIONTECH	Resident was found deceased in his bed at 7:15 am.
956903-1	MODERNA	mi Narrative: patient with asymptomatic covid 19, covid positive 12/10/2020.
956962-1	UNKNOWN MANUFACTURER	COVID 19 vaccine, unknown which company Chronically ill in a skilled nursing facility found diaphoretic, hypotensive, hypoxia to 85% arrived to Emergency dept in cardiac arrest Die
956966-1	MODERNA	hypoxia, secretions,cough, dyspnea Narrative: ALS patient on hospice with ongoing history of aspiration pna, receiving tube feeds. Developed incr in secretions, hypoxeia, temp and
956994-1	MODERNA	The patient had severe shortness of breath resulting in cardiac arrest on the 5th day after the vaccine. Shortness of breath started 12 hours after injection. On the 5th day, the patie
957116-1	MODERNA	Sudden death without warning symptoms 4 days after vaccine. Many medical problems which most likely explain the outcome but spouse feels it is related and it is a new vaccine. M
957163-1	PFIZER\BIONTECH	Resident received 1st on 1/11/21 at 12:10am (1/12/21) resident was found unresponsive. Code Blue, 911 called at 12:11am. FD and EMS arrived, resident pronounced at 12:51am.
957799-1	MODERNA	Presented to Urgent Care for weakness and confusion, transferred to ED, patient had a cardiac arrest and was unable to be resuscitated
958069-1	MODERNA	Started with cough, mild shortness of breath and feeling terrible in evening of 1/19.
958072-1	PFIZER\BIONTECH	Death 3 days after receiving 2nd dose of COVID vaccine, unknown if related to vaccine administration.
958228-1	MODERNA	Patient has end stage renal disease and rapidly worsening dementia, family could no longer care for him at home, and he was admitted for 14-day quarantine prior to admission to ir CREAT 6.93 K 5.2 were his baseline. He was found to be deceased on 1/18 at 11:18 pm.
958322-1	PFIZER\BIONTECH	Shaking and then became unresponsive
958443-1	PFIZER\BIONTECH	death by suicide Narrative: death by suicide; 12/26/20, self inflicted gun shot wound; found deceased by family member
958745-1	MODERNA	Resident was noted to have increase weakness on 1/15/2021. Resident was warm to touch with low grade fever of 99.3 F. Resident was up propelling self in w/c on 1/16/2021 he wa
958914-1	PFIZER\BIONTECH	Death on 1/15/2020
958935-1	MODERNA	Sudden Death within 24 hours of vaccine
958971-1	MODERNA	Hemorrhagic Stroke, Right Basal Ganglion
959001-1	MODERNA	Patient woke apx 0200 complaining of nausea to group home staff. Vitals were checked at that time and WNL. Patient went back to bed. When staff went to wake patient apx 0530,
959079-1	PFIZER\BIONTECH	On 1/9/2021 observed with elevated respirations of 38-42 per minute, BP manually 72/50. pulse is jumping rapidly between 110-16 bpm. oxygen sat 76% RA, resident refusing oxy Received order for morphine 2mg per hr as needed for elevated respirations and pain. Dr. also gave orders to D/C Tamsulosin and finasteride. Resident continue with decreased O2 s
959147-1	PFIZER\BIONTECH	Unknown as to any correlation with vaccine as this was a hospice patient that was already experiencing decline. Patient became Jaundice for approximately one week prior to expirin
959167-1	MODERNA	Patient received COVID 19 vaccine 01/14/2021. Patient died in his sleep 01/16/2021.
959179-1	PFIZER\BIONTECH	Patient received COVID-19 vaccination on 1/14/2021. On 1/17/2021, patient was transferred to Hospital s/p multiple cardiac arrests. Patient was hyperkalemic and in acute renal fai EF of 40-45% and elevated troponins. Patient was made DNR and placed on comfort care. Patient passed away on 1/18/2021. Ultimately we suspect that the patients condition was
959272-1	MODERNA	Patient died 4 days after immunization. Probably unrelated to immunization, as patient has been in poor health and was receiving hospice services. I have no details related to his ill
959356-1	MODERNA	Pt passed away the day after the vaccine was given.
959568-1	MODERNA	Patient received her first dose of the Moderna COVID-19 Vaccination on Saturday January 16th 2021 at approximately 12pm. She completed all necessary screening forms and was c CDPHE/CDC guidelines and left the Clinic in stable condition after her observation period was complete. On the morning of Tuesday, January 19th, 2021, the patient was found uncor transferred to other Hospital for higher level care. She was seen by neurosurgery and diagnosed with a ruptured aneurysm. She was treated in the ICU for 24 hours, at which point h
959591-1	MODERNA	Resident has increase weakness and lethargy with abnormal labs. He was transferred to the ER. He was admitted to the hospital and treated for worsening AKI and hypotension.
959729-1	PFIZER\BIONTECH	Per Nursing Staff- patient died within 24 hours of receiving the vaccine. patient has hospice. Please contact director of nursing for more details.
959747-1	PFIZER\BIONTECH	per staff at facility patient died 24 hours post vaccination. Please contact Director of Nursing for further details.
959929-1	PFIZER\BIONTECH	"Narrative: Patient seen in ED 1-17-21 with c/c of ""bloated with epigastric pain"". Patient with complicated medical history including stage 1B pancreatic cancer (was currently on ch history: CAD s/p CABG 2009, PAF, and HTN). Regarding ER visit for epigastric pain, nothing notable was found on workup and patient was to discharge home to rest. There were ava notable issues. The following day, Monday 1-18-21, patient's caregiver called facility at 22:30 to report he had a fever of 102.8 degrees and that he had been ""feeling kind of bad al myalgia. WBC WNL, CXR unremarkable for infection, UA neg for bacteria, LFTs WNL, blood cultures negative. Procalcitonin elevated at 17.8 -- suggesting inflammatory response. Pat Patient then coded and resuscitation was attempted for approximately 30 minutes. Patient did not survive the code. Coroner has been notified and family is considering autopsy at ti
960426-1	PFIZER\BIONTECH	expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar death events for 8 patients. This report is for 1st of 8 pa covid-19 immunisation. The patient medical history was and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details espec safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in res different patient;US-PFIZER INC-2021034598 same drug, reporter and event but different patient;US-PFIZER INC-2021034599 same drug, reporter and event but different patient;U same drug, reporter and event but different patient;US-PFIZER INC-2021034596 same drug, reporter and event but different patient.; Reported Cause(s) of Death: expired before r
960427-1	PFIZER\BIONTECH	expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 2nd of 8 patients covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details especially evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response,
960428-1	PFIZER\BIONTECH	expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 3rd of 8 patients covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. It completely exclude the possible causality between the reported death and the administration of COVID 19 vaccine, bnt162b2. However, more information on the patient's underlying impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate dat Death: expired before receiving the second dose
960429-1	PFIZER\BIONTECH	expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 4th of 8 patient. covid-19 immunization. The patient's medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date. T very limited for full assessment. The patient died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, co evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this revie 2021034595 same drug, reporter and event but different patient; Reported Cause(s) of Death: expired before receiving the second dose
960430-1	PFIZER\BIONTECH	7 residents expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar events for 8 patients. This report is for 6th c SINGLE DOSE for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient expired before receiving the second dose on an BNT162b2 vaccine and documented as such in the global safety database until sufficient information is available to allow an unrelated causality assessment. The impact of this report concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as ap
960431-1	PFIZER\BIONTECH	expired before receiving the second dose; This is a spontaneous report from a contactable nurse. This nurse reported similar death events for 8 patients. This report is for 7th of 8 pa covid-19 immunisation. The patient medical history was and concomitant medications were not reported. The patient expired before receiving the second dose on an unspecified date died following the vaccine use; further information such as patient demographics, complete medical history, concomitant medications, concurrent illness and event term details espec safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in res different patient; Reported Cause(s) of Death: expired before receiving the second dose
960437-1	PFIZER\BIONTECH	platelets dropped so low/thrombocytopenia; Hemorrhagic stroke/brain hemorrhage; This is a spontaneous report from a contactable nurse. A 56-year-old male patient received BNT were unknown. The reporter read about the doctor that died that developed thrombocytopenia after taking the vaccine, stated it was in the news yesterday. The patient received the had specialists that tried to get his platelet count back up again and they could not get his platelets back up again and he ended up having the hemorrhagic stroke. The reporter alre Comments: Very limited information is currently available. Lacking patient's underlying medical conditions, clinical course, relevant lab data, the Company cannot make a meaningfu the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. An stroke/brain hemorrhage; platelets dropped so low/thrombocytopenia

VAERS ID	Vaccine Manufacturer	Adverse
960460-1	PFIZER\BIONTECH	"died; tested positive for COVID; tested positive for COVID; This is a spontaneous report from a contactable consumer from a Pfizer-sponsored program, Pfizer First Connect. A 97-y dose for COVID-19 immunization; administered by the nursing home. Medical history included glaucoma from an unknown date and unknown if ongoing. Concomitant medications in significant). The patient died (death, medically significant) on 17Jan2021. The clinical course was reported as follows: The reporter stated that in regard to the patient's height and w pounds. The reporter stated that her father was in a nursing home. The patient received his first dose of the COVID vaccine on 30Dec2020. The patient died on 17Jan2021. The repo The reporter stated that there was nothing wrong with her dad. He was elderly with no health issues. ""He was literally on no medications. The only reason he was in the nursing hor after the vaccine, the patient tested positive for COVID ""like all the other people"" (no further details provided). The reporter stated that her dad had no symptoms of COVID. The d again stated that the patient literally had nothing wrong with him. ""They were shocked. They fed him and he took a nap. He was sleeping, but it was eternally."" The reporter stated and procedures which included COVID-19 virus test: positive on 07Jan2021. History of all previous immunization with the Pfizer vaccine considered as suspect: none. It was unknow events following the prior vaccinations. The clinical outcome of the event, died, was fatal. The clinical outcome of the event, tested positive for COVID, was unknown. The patient die will be requested during follow up.; Reported Cause(s) of Death: died"
960552-1	PFIZER\BIONTECH	At approximately 930am I arrived at Memory Care. I met with the director of the facility and she directed me to where my team would be setting up. My team consisted of (technicia minutes and maybe even longer . I also explained that we would need one of her staff monitoring while we vaccinate. She agreed, and proceeded to designate her staff and the cafe also stop occasionally to mix the vaccine and check the temperature of the aero safe. At approximately 12:50pm, the director rushed in and stated that a patient is not responding, a Once we got to the room, the patient was in bed and there were 4 staff members standing bedside and one of them turned and stated the patient has passed. At that point I asked t detiorating and had not eaten or drank anything all day . They also stated that the patient had been monitored for 15 minutes post vaccination. I then left the room and grabbed the administration information for Immunizer Section was filled out by Nurse. I then proceeded to ask the director once again if there were staff that was monitoring her for 15 minutes, explained the situation. After speaking to corporate, I also asked nurse, if she remembered the patient. She stated that she did and at the time of the vaccination the patient was no how she usually is and if its ok to vaccinate her. Both Staff members stated that it its ok,this is how she is. The Nurse then proceeded to vaccinate. At approximately 3:10pm, as I wa stated that on Friday, Jan 15th,2021, they had informed the family that the patient was rapidly detiorating. Staff also stated that the family knowingly gave the consent to vaccinate the Vaccine. Per Lead Pharmacist at the clinic.
960752-1	MODERNA	Extreme Fatigue
960841-1	PFIZER\BIONTECH	Patient developed 104.4 temp approximately 48 hours after being given the vaccine. I treated him with antibiotics, IV fluids, cooling methods. CXR does show a new right perihilar in
961010-1	PFIZER\BIONTECH	Resident returned to the memory support unit at 1500. Resident was than toileted and transferred in to bed per his request. At 1515 resident was observed face down beside bed, re EMS.
961339-1	MODERNA	possibly got it at clinic, possibly who administered shot. Pts. daughter said the pts boyfriend denied any symptoms the whole day but that in the middle of the night the pt passed av
961434-1	PFIZER\BIONTECH	This is a 94-year-old male who is brought in by ambulance after being found on the floor with unknown downtime. He was in asystole upon EMS arrival. He remains in asystole. No a vaccination this morning. The patient is evaluated emergently. CPR was ongoing with 3 rounds of epinephrine given. The patient remains in asystole. He has rigor mortis. The patient patient's time of death is 2113.
961705-1	PFIZER\BIONTECH	approximately 3 hours prior to expiring the patient was experiencing forceful emesis. later was found to have expired, patient was comfort care only.
961741-1	PFIZER\BIONTECH	Theápatientáreceivedáhisávaccineáináthe.ámorningáofá1/20/2021,áwhileágettingáintoácarátoágoáseeáhisápulmonologist,áaboutá2áhoursáafter,ácollapsed,áunresponsiveáwithháasys
961776-1	PFIZER\BIONTECH	1/13/2021 12:00 PM: Patient received COVID-19 Vaccine. 1/14/2021 21:00: Nurse performed routine rounds and the patient appeared okay. 1/14/2021 22:00: CNA discovered pati
961845-1	MODERNA	Narrative:
961848-1	PFIZER\BIONTECH	"Narrative: Was pt previously covid positive?- Yes. Initial- 10/27/2020, 11/29/2020, 12/22/2020 Are there any predisposing factors for patient experiencing adverse drug event?- Ye Atorvastatin, Mirtazapine and Duloxetine. Pt with 2 doses of covid-19 vaccine, second one on 01/08/2021, 2 days pre-death Any occurrence of an ADR at time of administration? Did 12/21/2021, LFTS increased ~01/01/2021, peaked on 01/03/2021 and were decreasing on 01/07/2021 Was there an ADR between observation period and date of death? No Did pa death--was hospitalization attributable to ADE? Yes re-admitted to inpatient on 12/31/2020. GI bleed Is there an alternative cause of death? Yes, as noted above. Quite a complicate
962307-1	PFIZER\BIONTECH	tired; legs felt heavy; stopped breathing; This is a spontaneous report from a Pfizer-sponsored program a non-contactable consumer. A 93-year-old male patient received bnt162b2 medications were not reported. Patient received vaccine around 11:00 a.m. About two hours later, he said he was tired and couldn't continue with the physical therapy he was doing. reported if an autopsy was performed. Outcome of stopped breathing was fatal. Outcome of tired and legs felt heavy was unknown. No follow-up attempts are possible; information
962308-1	PFIZER\BIONTECH	died; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that an 83-year-old female patient (reporter mother) received BNT162B2 (PFI hospice care and dementia. The patient's concomitant medications were not reported. The patient died one day after getting vaccine. She was reportedly in good health the day befo stable in her health. The reporter read investigating 23 deaths of people receiving vaccine in similar conditions. The patient died on an unspecified date. It was not reported if an aut
962318-1	MODERNA	"Called to schedule second vaccine and daughter reports that he died on01/19/2021 with ""COVID"""
962325-1	PFIZER\BIONTECH	"Patient's wife called this morning stating that her husband has passed away last night. After receiving first dose of Pfizer COVID-19 vaccine at around 0830, patient remained in the ""dissipated after a few minutes"" followed by a headache which ""dissipated after a few minutes"" as well. Then patient complained of nausea, no vomiting and ""couldn't relax."" Pe around 2100/2200, patient was quiet and when she checked on him, ""he wasn't responding anymore."" Wife then called 911, ""but they couldn't revive him."""
962390-1	PFIZER\BIONTECH	Admitted to hospital after vaccination with Acute hypoxemic respiratory failure, Septic shock; Aneurysm of arteriovenous dialysis fistula; expired 1/16/2021
962714-1	PFIZER\BIONTECH	We do not believe that the patient's death was an adverse event from the vaccine. Patient received COVID vaccine from Pfizer Dose #1 12/19/2020 (lot # EK5730) and Dose #2 1/7 congestive heart failure after recent aspiration pneumonia requiring hospitalization. Death was anticipated and not sudden. We were told to report his death to VAERS even though h
962716-1	PFIZER\BIONTECH	Patient deceased
962764-1	PFIZER\BIONTECH	Patient did not have any adverse reaction to the COVID vaccine, but we were asked by our health dept to submit a VAERS report since the patient died between his first and second pneumonia complicated by end-stage heart failure and ischemic cardiomyopathy. Death was anticipated and not sudden.
962784-1	PFIZER\BIONTECH	patient expired 1/15/2021; had been treated as outpatient for pneumonia, likely COVID-19 but no positive test result in December 2020. PMH diabetes
962827-1	PFIZER\BIONTECH	Admitted 1/14/21: Patient is an elderly 93-year-old female with multiple medical problems including chronic combined CHF, P 80, diabetes mellitus, HTN, hyperlipidemia, CKD stage the hospital, condition clinically deteriorated. Patient elected to be DNR/DNI while in the ED. Patient was pronounced dead at 10:30 p.m. earlier today. Preliminary cause of death: H
962940-1	MODERNA	Pt received second dose of COVID vaccine on 01/20/2021 at 1430. At 1600 Pt developed a wet productive cough with coarse crackles. Pt ate dinner at 5 pm cough persisted. At 18:3 sitting in the lounge and a CNA noticed that Pt was pale/white in color and clammy. 02 Sat was 85%. Respirations were labored. Pt was placed on 4 L of 02. Increased to 5 L via face
962966-1	PFIZER\BIONTECH	On Saturday, 1/16/2021, Patient went to the grocery store. Upon her return, she indicated she was experiencing N/V and some throat swelling. Patient subsequently collapsed and e are pending and the Coroners investigation is ongoing. Spouse believes that her death was caused by the vaccine.
962995-1	MODERNA	No immediate reaction. Patient-reported deceased four days later on Jan. 19, 2021. As of this date cause of death is unknown to our clinic.
963016-1	MODERNA	unknown. Event occurred after leaving vaccination site
963057-1	PFIZER\BIONTECH	presented to ED 1/9/21 with abdominal pain, progressive worsening weakness and fatigue and new onset A fib with RVR likely due to hypertensive urgency . Patient progressed clini clinical decline, palliative care had been consulted on end of life care. Patient expired 1/17/21
963163-1	MODERNA	Narrative:
963167-1	MODERNA	Narrative: Symptoms: & Cardiac Arrest; Death Treatment: EPINEPHRINE
963235-1	MODERNA	Patient diagnosed with COVID on January 9, 2021 after being exposed to family member that was under quarantine in the same household. Admitted to the hospital and was dischar
963269-1	MODERNA	Patient passed away on 01/18/2021
963388-1	MODERNA	Patient died unexpectedly 5 days after receiving vaccine (1/10/2021).
963610-1	MODERNA	Patient deceased on 01/17/2021
963902-1	PFIZER\BIONTECH	Death; This is a spontaneous report from four non-contactable consumers via a Pfizer-sponsored program Corporate (Pfizer) Social Media Platforms. A 78-year-old male patient recei included Alzheimer's Disease, encephalopathy, hypertension, acute kidney failure, urinary retention and recent urinary tract infection (UTI), all from an unspecified date. Concomitan hydrocodone bitartrate, paracetamol (HYDROCODONE/ACETAMINOPHEN), loperamide (MANUFACTURER UNKNOWN), ondansetron (MANUFACTURER UNKNOWN), senna alexandrina (adverse events and no issues on 29Dec2020. The patient died on 30Dec2020, at approximately 2:00 AM. It was unknown if an autopsy was performed. It was unknown if the event Death: Death
964401-1	MODERNA	Pt died 4 days after vaccine, no known reaction to the vaccination
964617-1	PFIZER\BIONTECH	Death, which I believe is unrelated to vaccination
964629-1	PFIZER\BIONTECH	Death - Hospice patient with metastatic CA admitted to facility and received vaccine during stay. No adverse sequelae noted from vaccine administration, but reporting as required b administered by protocol without complications. The patient had been asked and denied any prior severe reaction to this vaccine or its components and gave permission to receive it to underlying terminal illness.
964636-1	PFIZER\BIONTECH	Pt on hospice in facility for severe cardiomyopathy unable to perform interventions received vaccine without adverse sequelae died 5 days later. Reporting as required. Narrative: Rep following the immunization related to the vaccine. The patient denied any prior severe reaction to this vaccine or its components, and the patient gave verbal consent to receive the with very poor potential to heal due to advanced PVD.
964653-1	PFIZER\BIONTECH	loss of consciousness; respiratory distress Narrative: Patient tolerated his 1st dose of the COVID-19 vaccine well, on 12/16/2020, and received his 2nd dose on 1/6/2021. Patient ha on the evening of 1/6/21, which was effectively managed, but by early morning he spiked a fever of 102.9 with a sat of 86.1%. He continued to deteriorate from that point on and d
964671-1	PFIZER\BIONTECH	Death on 1-5-21
964724-1	PFIZER\BIONTECH	Death 1-15-21

Note: Submitting a report to VAERS does not mean that healthcare personnel or the vaccine caused or contributed to the adverse event (possible side effect).

Notes:

Caveats: VAERS accepts reports of adverse events and reactions that occur following vaccination. Healthcare providers, vaccine manufacturers, and the public can submit reports to VAERS. While very important in monitoring vaccine safety, VAERS reports alone cannot be used to determine if a vaccine caused or contributed to an adverse event or illness. The reports may contain information that is incomplete, inaccurate, coincidental, or unverifiable. Most reports to VAERS are voluntary, which means they are subject to biases. This creates specific limitations on how the data can be used scientifically. Data from VAERS reports should always be interpreted with these limitations in mind.

The strengths of VAERS are that it is national in scope and can quickly provide an early warning of a safety problem with a vaccine. As part of CDC and FDA's multi-system approach to post-licensure vaccine safety monitoring, VAERS is designed to rapidly detect unusual or unexpected patterns of adverse events, also known as "safety signals." If a safety signal is found in VAERS, further studies can be done in safety systems such as the CDC's Vaccine Safety Datalink (VSD) or the Clinical Immunization Safety Assessment (CISA) project. These systems do not have the same limitations as VAERS, and can better assess health risks and possible connections between adverse events and a vaccine.

Key considerations and limitations of VAERS data:

- Vaccine providers are encouraged to report any clinically significant health problem following vaccination to VAERS, whether or not they believe the vaccine was the cause.
- Reports may include incomplete, inaccurate, coincidental and unverified information.
- The number of reports alone cannot be interpreted or used to reach conclusions about the existence, severity, frequency, or rates of problems associated with vaccines.
- VAERS data are limited to vaccine adverse event reports received between 1990 and the most recent date for which data are available.
- VAERS data do not represent all known safety information for a vaccine and should be interpreted in the context of other scientific information.

Some items may have more than 1 occurrence in any single event report, such as Symptoms, Vaccine Products, Manufacturers, and Event Categories. If data are grouped by any of these items, then the number in the Events Reported column may exceed the total number of unique events. If percentages are shown, then the associated percentage of total unique event reports will exceed 100% in such cases. For example, the number of Symptoms mentioned is likely to exceed the number of events reported, because many reports include more than 1 Symptom. When more then 1 Symptom occurs in a single report, then the percentage of Symptoms to unique events is more than 100%. More information. (/wonder/help/vaers.html#Suppress)

Data contains VAERS reports processed as of the previous Friday. The VAERS data in WONDER are updated weekly, yet the VAERS system receives continuous updates including revisions and new reports for preceding time periods. More information. (/wonder/help/vaers.html#Reporting)

Values of Event Category field vary in their availability over time due to changes in the reporting form. The "Emergency Room/Office Visit" value was avaliable only for events reported using the VAERS-1 form, active 07/01/1990 to 06/29/2017. The "Congenital Anomaly/Birth Defect", "Emergency Room", and "Office Visit" values are available only for events reported using the VAERS 2.0 form, active 06/30/2017 to present. These changes must be considered when evaluating count of events for these categories.

Help: See The Vaccine Adverse Event Reporting System (VAERS) Documentation (/wonder/help/vaers.html) for more information.

Query Date: Jan 29, 2021 8:56:57 PM

Suggested Citation:

United States Department of Health and Human Services (DHHS), Public Health Service (PHS), Centers for Disease Control (CDC) / Food and Drug Administration (FDA), Vaccine Adverse Event Reporting System (VAERS) 1990 - Previous Friday, CDC WONDER On-line Database. Accessed at http://wonder.cdc.gov/vaers.html on Jan 29, 2021 8:56:57 PM

Query Criteria:

Event Category:	Death
State / Territory:	The United States/Territories/Unknown
Vaccine Products:	COVID19 VACCINE (COVID19)
VAERS ID:	All
Group By:	VAERS ID; Vaccine Manufacturer
Show Totals:	False
Show Zero Values:	Disabled